

Food and Drug Administration Rockville, MD 20857

August 25, 2004

FILE COPY

Anthony C. Celeste, President AAC Consulting Group Suite 500 Rockville, MD 20855-2765

Dear Mr. Celeste:

Your petition requesting the Food and Drug Administration to determine whether Penthrane (methoxyflurane, liquid, 99.9%) inhalation (NDA 13-056), manufactured by Abbott Laboratories, has been voluntarily withdrawn from sale for safety or efficacy reasons was received by this office on 08/25/2004. It was assigned docket number 2004P-0379/CP1 and it was filed on 08/25/2004. Please refer to this docket number in future correspondence on this subject with the Agency.

Please note that the acceptance of the petition for filing is a procedural matter in that it in no way reflects an agency decision on the substantive merits of the petition.

Sincerely,

Lyle LY Jaffe

Division of Dockets Management Office of Management Programs

Office of Management

2004P-0379

ACKI